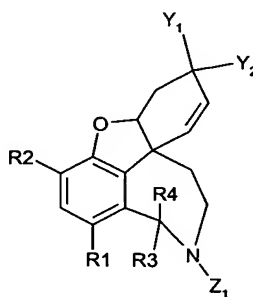


## Patent claims:

1. Use of galanthamine and galanthamine derivatives exhibiting cholinergic activity for manufacturing medicaments for the treatment of post-operative delirium and/or subsyndromes of post-operative delirium.

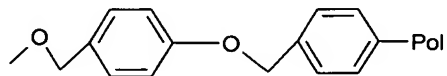
2. Use according to claim 1 for manufacturing medicaments for the preventive treatment of post-operative delirium and/or subsyndromes of post-operative delirium.

3. Use according to claims 1 or 2, characterized by the fact that the galanthamine derivatives have the general formula



Ia

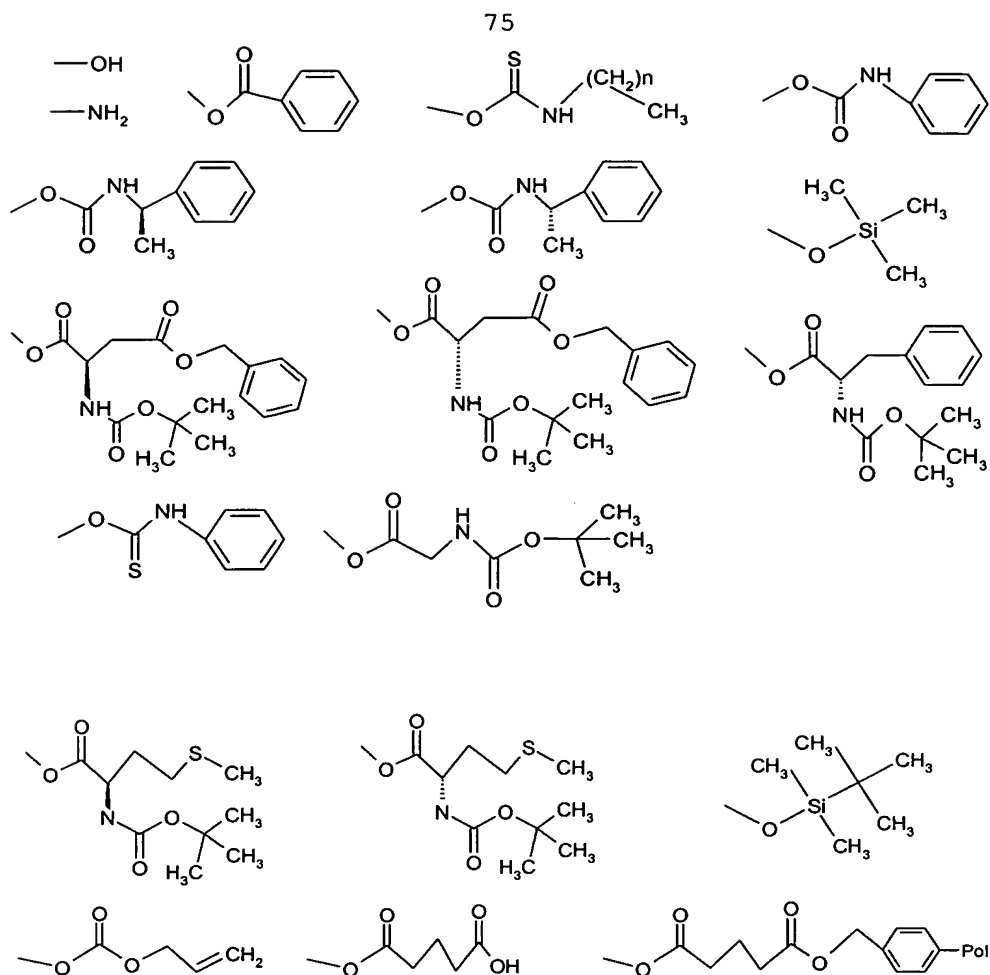
and the salts thereof, wherein  $R_1$  is H, branched or straight chain ( $C_1$ - $C_6$ ) alkyl, Br,  $NO_2$ ,  $NR_5R_6$  wherein  $R_5$  and  $R_6$  are the same or different and are selected from H, branched or straight chain ( $C_1$ - $C_6$ ) alkyl, and wherein  $R_2$  is OH, branched or straight chain ( $C_1$ - $C_6$ ) alkyl, methoxy, phenyloxy or the following group



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whereby Pol is a polymer, preferably one in accordance with WO-A1-01/174820, and wherein  $R_3$  and  $R_4$  either at the same time or alternatively are H, D, CN, straight chain or branched ( $C_1$ - $C_6$ ) alkyl or a carbonyl group together, wherein  $Y_1$  and  $Y_2$  alternatively are H or a group selected from:

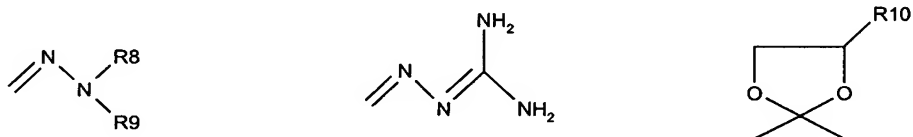
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wherein n represents a value of 0, 1 to 15, and Pol has the meaning indicated above, and wherein Y<sub>1</sub> and Y<sub>2</sub> further represent together a carbonyl group (=O), =NH, = N-OR<sub>7</sub>, wherein R<sub>7</sub> is H, tosylate or branched or straight chain (C<sub>1</sub>-C<sub>6</sub>) alkyl, or Y<sub>1</sub> and Y<sub>2</sub> together is a group selected from:

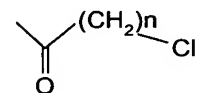
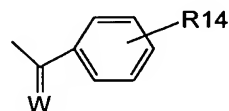
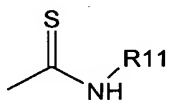
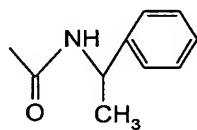
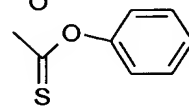
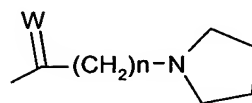
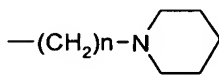
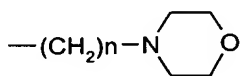
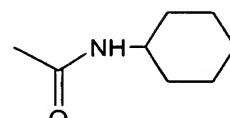
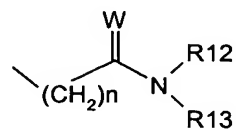
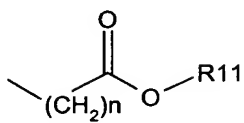
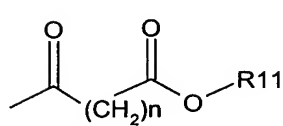
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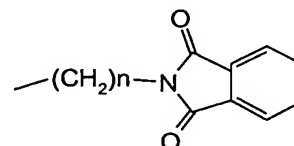
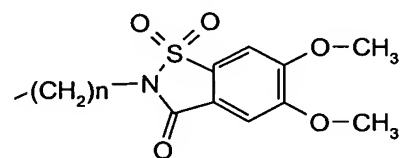
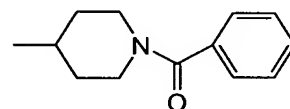
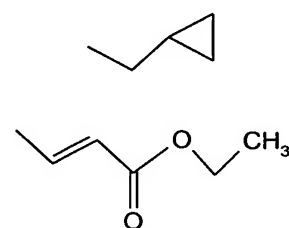
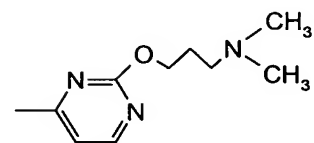
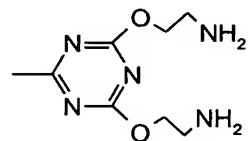


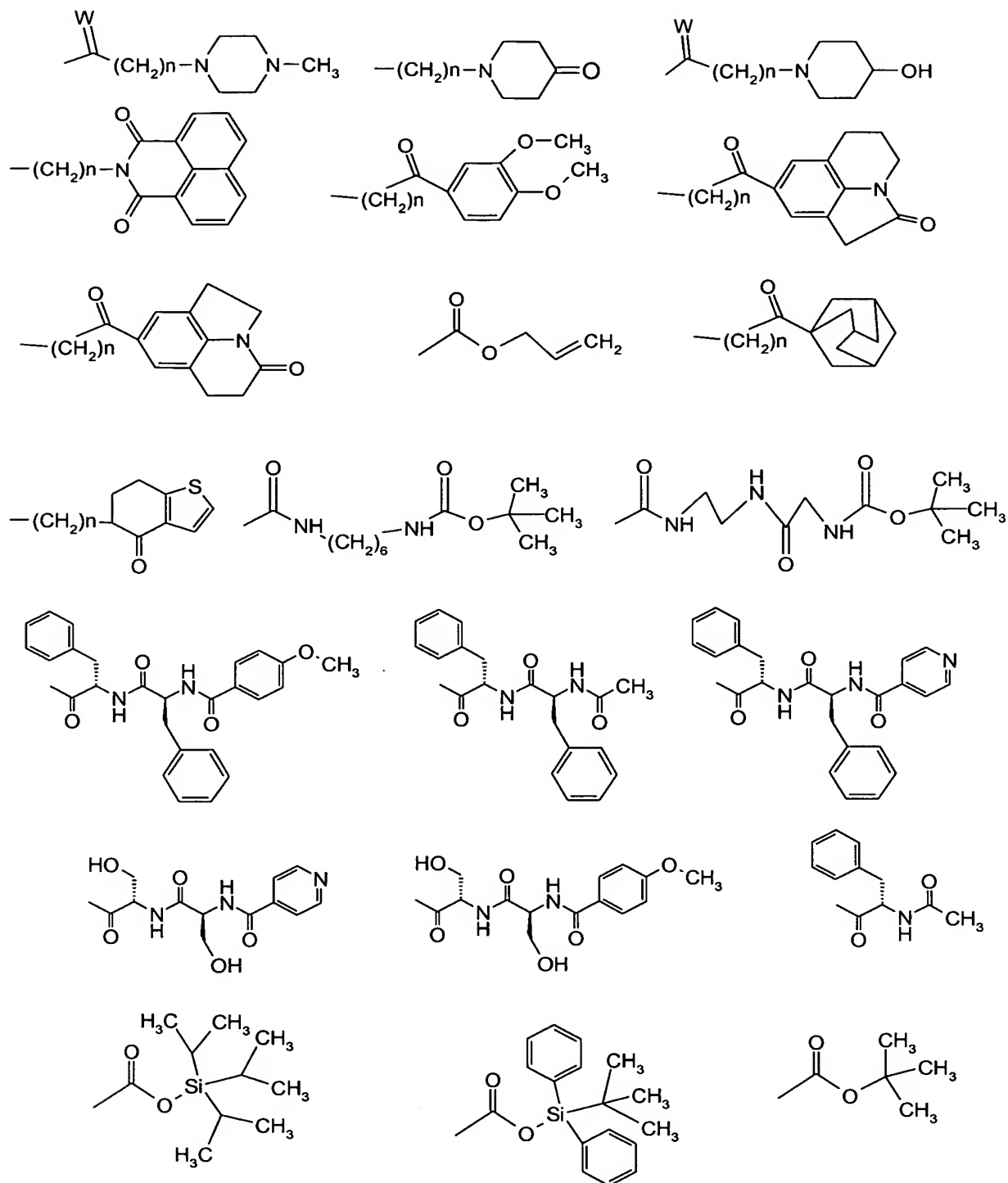
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wherein R<sub>8</sub> and R<sub>9</sub> are the same or different and are H, branched or straight chain (C<sub>1</sub>-C<sub>6</sub>) alkyl, -(CH<sub>2</sub>)<sub>2</sub>-OH, CHO, CONH<sub>2</sub>, tBOC (tert-Butoxycarbonyl), or mean -COCOOH, R<sub>10</sub> is H or CH<sub>3</sub>, and wherein when Y<sub>1</sub> is -O-(CH<sub>2</sub>)<sub>2</sub>-OH, Y<sub>2</sub> is OH, and wherein Z<sub>1</sub> is H, branched or

straight chain (C<sub>1</sub>-C<sub>6</sub>) alkyl, (C<sub>2</sub>-C<sub>7</sub>) alkenyl, (C<sub>2</sub>-C<sub>7</sub>) alkynyl, tri-fluoroacetyl, formyl, phenyl or a group selected from:



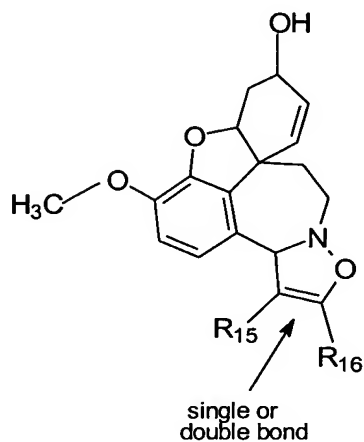




wherein  $R_{11}$  is H, straight chain ( $C_1$ - $C_6$ ) alkyl, branched ( $C_1$ - $C_6$ ) alkyl or ( $C_2$ - $C_7$ ) alkenyl,  $R_{12}$  and  $R_{13}$  are the same or different and are selected from H, straight chain or branched ( $C_1$ - $C_6$ ) alkyl,

phenyl, chlorophenyl, (trifluoromethyl)-phenyl or 1-naphtyl, wherein  $R_{14}$  is H, F,  $CH_3$ ,  $NO_2$ , Cl, Br, J,  $CF_3$ , n has the meaning indicated above, m is 0 or 1, and W has the meaning H or O, and wherein further  $Z_1$  and  $R_3$  form a common ring

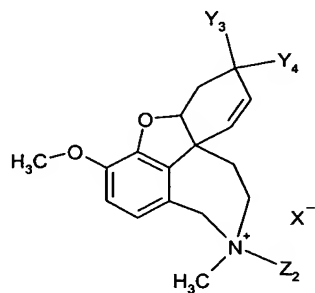
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wherein  $R_{15}$  and  $R_{16}$  alternatively mean H,  $COOCH_3$ ,  $COOCH_2CH_3$ , CN,  $COCH_3$ .

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4. Use according to claims 1 or 2, characterized by the fact that the used Galanthamine derivatives have the general formula Ib



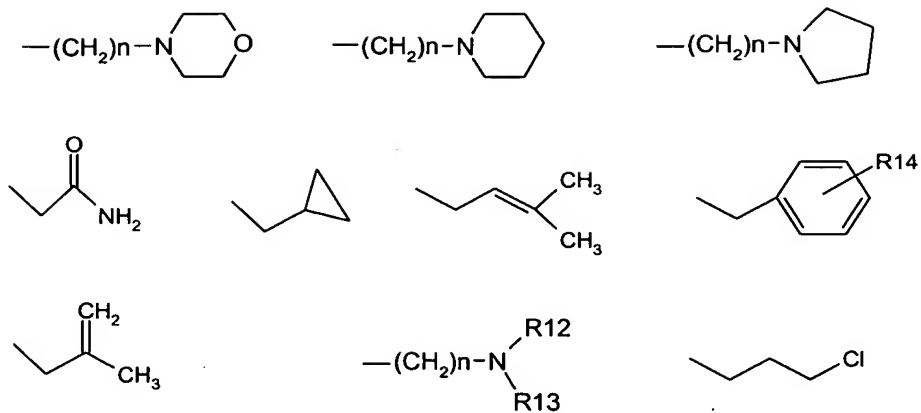
Ib

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wherein  $Y_3$  and  $Y_4$  alternatively mean H and OH, X is Cl, Br or I,  $Z_2$  is oxygen (N-oxide and no counterion), branched or straight chain ( $C_1-C_6$ ) alkyl, or ( $C_2-C_7$ ) alkenyl or ( $C_2-C_7$ ) alkynyl or a group selected from:

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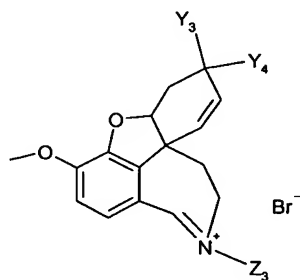
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wherein  $n$ ,  $R_{12}$ ,  $R_{13}$  and  $R_{14}$  have the meanings as defined in accordance with claim 3.

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5. Use according to claims 1 or 2, characterized by the fact that the used galanthamine derivatives have the general formula Ic



Ic

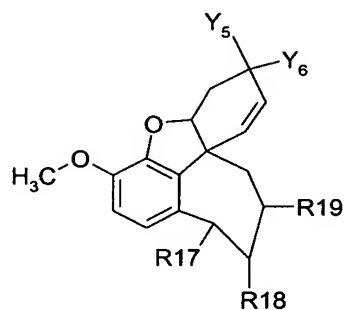
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wherein  $Y_3$  and  $Y_4$  the meaning defined in accordance with claims 3 or 4 have, and  $Z_3$  is oxygen (N-oxide and no counterion) or is a methyl.

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6. Use according to claims 1 or 2, characterized by the fact

that the used galanthamine derivatives have the general formula Id

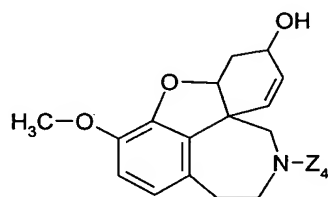


Id

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and their salts, wherein  $Y_5$  and  $Y_6$  alternatively are H or OH, or together form a keto group, and  $R_{17}$ ,  $R_{18}$ ,  $R_{19}$  are alternatively for two substituents H, wherein the third substituent is  $NH_2$  or  $CONH_2$ .

10 7. Use according to claim 1 or 2, characterized by the fact that the used galanthamine derivatives have the general formula Ie

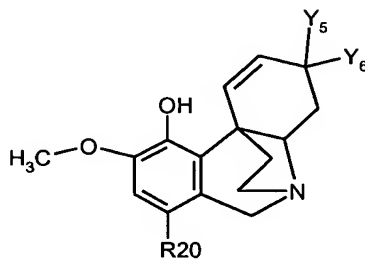


Ie

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or their salts, wherein  $Z_4$  is straight chain or branched ( $C_1-C_6$ ) alkyl or 4-brombenzyl.

20 8. Use according to claims 1 or 2, characterized by the fact that the used galanthamine derivatives have the general formula If:

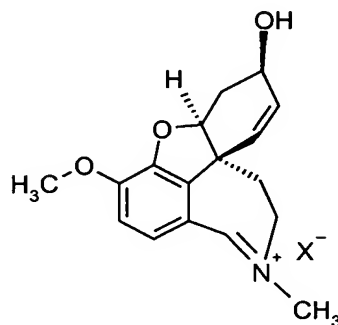


If



or their salts, wherein  $Y_5$  and  $Y_6$  have the meanings as defined in claims 3 to 7, and  $R_{20}$  is H or Br.

- 5           9. Use according to claims 1 or 2, characterized by the fact that the used galanthamine derivative has the following structural formula



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- and its pharmaceutical acceptable salts, hydrate or a solvate thereof and having the chemical name (4aS, 6R, 8aS)-6-Hydroxy-3-methoxy-11-methyl-4a,5,9,10-tetrahydro-6H-benzofuro[3a,3,2-f][2]benzazepinium.

10. Use according to claim 9, characterized by the fact that the pharmaceutical acceptable salt counterion of (4aS, 6R, 8aS)-6-Hydroxy-3-methoxy-11-methyl-4a,5,9,10-tetrahydro-6H-benzofuro[3a,3,2-ef][2]benzazepinium is selected from the group of halides, preferably bromide, carboxylic acids with 1-3 carboxyl functions, particularly preferably tartrate, malonate, fumarate and succinate, and sulfonic acids, preferably methane sulfonic acid.